

Atp8b1 Cas9-CKO Strategy

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Design Date: 2020-1-17

Project Overview



Project Name

Atp8b1

Project type

Cas9-CKO

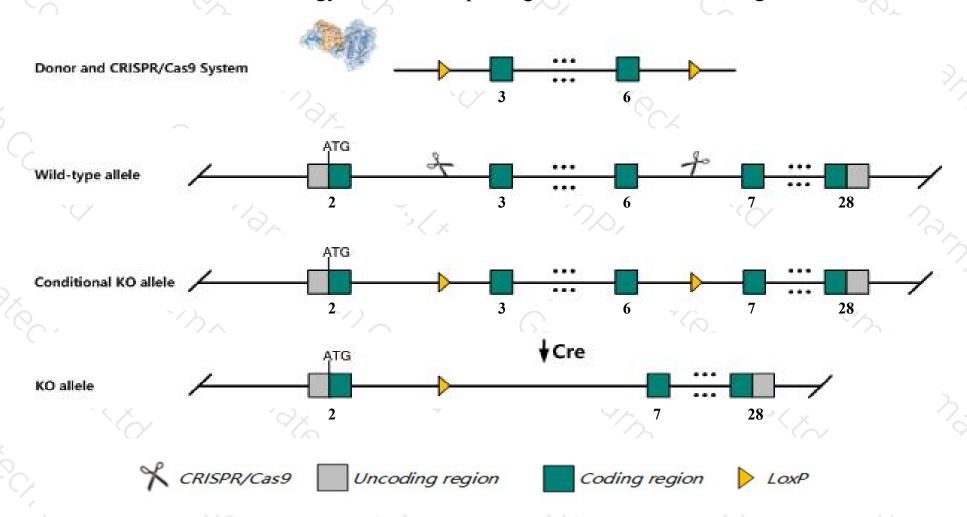
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Atp8b1 gene. The schematic diagram is as follows:



Technical routes



- The *Atp8b1* gene has 3 transcripts. According to the structure of *Atp8b1* gene, exon3-exon6 of *Atp8b1-201* (ENSMUST00000025482.9) transcript is recommended as the knockout region. The region contains 373bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Atp8b1* gene. The brief process is as follows:CRISPR/Cas9 system and Donor were microinjected into the fertilized eggs of C57BL/6JGpt mice. Fertilized eggs were transplanted to obtain positive F0 mice which were confirmed by PCR and sequencing. A stable F1 generation mouse model was obtained by mating positive F0 generation mice with C57BL/6JGpt mice.
- The flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.

Notice



- According to the existing MGI data, Homozygous mice display abnormal bile salt homeostasis, normal bile secretion, and an impaired ability to handle increased bile salt loading resulting in liver damage and weight loss on a bile salt supplemented diet.
- Transcript 202 CDS 5' and 3' incomplete the influences is unknown.
- The *Atp8b1* gene is located on the Chr18. If the knockout mice are crossed with other mice strains to obtain double gene positive homozygous mouse offspring, please avoid the two genes on the same chromosome.
- ➤ This Strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risk of loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

Gene information (NCBI)



Atp8b1 ATPase, class I, type 8B, member 1 [Mus musculus (house mouse)]

Gene ID: 54670, updated on 31-Jan-2019

Summary

↑ ?

Official Symbol Atp8b1 provided by MGI

Official Full Name ATPase, class I, type 8B, member 1 provided by MGI

Primary source MGI:MGI:1859665

See related Ensembl: ENSMUSG00000039529

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as Al451886, FIC1, Ic

Expression Biased expression in large intestine adult (RPKM 22.0), colon adult (RPKM 19.6) and 11 other tissuesSee more

Orthologs <u>human</u> all

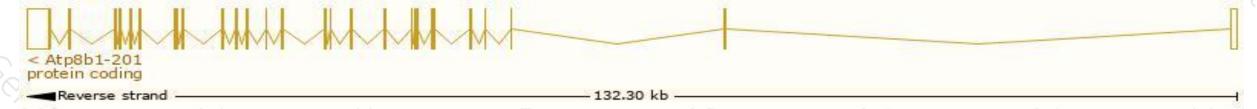
Transcript information (Ensembl)



The gene has 3 transcripts, all transcripts are shown below:

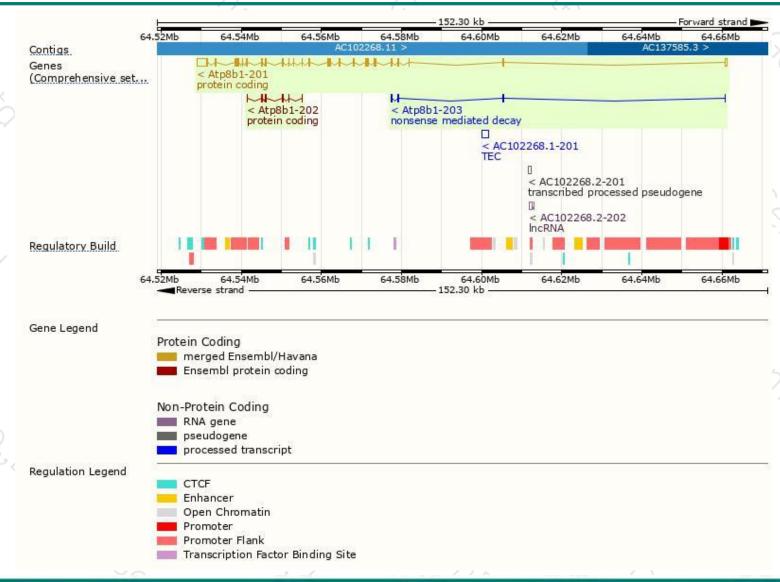
	- AMDE					2014a	
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Atp8b1-201	ENSMUST00000025482.9	6719	<u>1251aa</u>	Protein coding	CCDS29304	Q148W0	TSL:1 GENCODE basic APPRIS P1
Atp8b1-202	ENSMUST00000235459.1	737	245aa	Protein coding	+1	89	5' and 3' truncations in transcript evidence prevent annotation of the start and the end of the CDS. CDS 5' and 3' incomplete
Atp8b1-203	ENSMUST00000237686.1	531	61aa	Nonsense mediated decay	20	82	

The strategy is based on the design of Atp8b1-201 transcript, The transcription is shown below



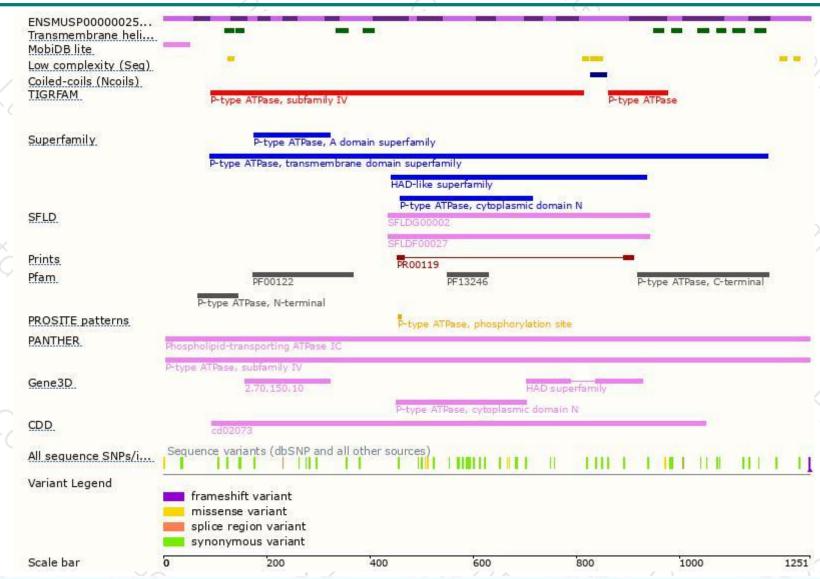
Genomic location distribution





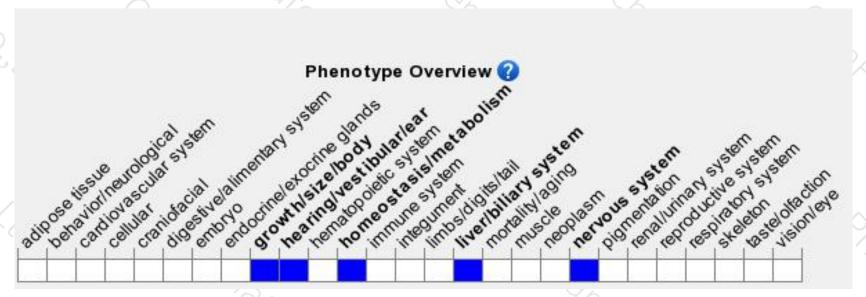
Protein domain





Mouse phenotype description(MGI)





Phenotypes affected by the gene are marked in blue.Data quoted from MGI database(http://www.informatics.jax.org/).

According to the existing MGI data, Homozygous mice display abnormal bile salt homeostasis, normal bile secretion, and an impaired ability to handle increased bile salt loading resulting in liver damage and weight loss on a bile salt supplemented diet.



If you have any questions, you are welcome to inquire. Tel: 400-9660890





